

In re Application of: Evans et al.  
Application No.: 09/155,252  
Filing Date: September 21, 1998  
Page 2 of 22

PATENT  
Attorney Docket No.: SALK1470-2  
(088802-1852)

-- Presently preferred response elements contain at least one copy (with one, two or three copies most common) of the minimal sequence:

AGGACA A AGGTCA (SEQ ID NO:5).

As noted above, the minimal sequence can optionally be flanked by additional residues, for example, as in the sequence:

GGACC AGGACA A AGGTCA CGTTC (SEQ ID NO:6).

Please replace the paragraph beginning at line 20 of page 15 with the following replacement paragraph:

-- Exemplary PPREs have been described in detail hereinabove.

Exemplary GAL4 response elements are those containing the palindromic 17-mer:

5' - CGGAGGACTGTCCTCCG - 3' (SEQ ID NO:7),

such as, for example, 17MX, as described by Webster et al., in Cell 52:169-178 (1988), as well as derivatives thereof. Additional examples of suitable response elements include those described by Hollenberg and Evans in Cell 55:899-906 (1988); or Webster et al. in Cell 54:199-207 (1988).

Please replace the paragraph beginning at line 30 of page 21 with the following replacement paragraph:

-- A basic vector useful for the generation of GAL4-receptor fusion proteins is called pCMX-GAL4 (see SEQ ID NO:3). This vector encodes GAL4 DNA binding domain, followed by a polylinker sequence useful in the cloning. The parental expression vector pCMX has been described by Umesono et al., in Cell 65:1255-1266 (1991), and the GAL4 portion of pCMX-GAL4 is derived

In re Application of: Evans et al.  
Application No.: 09/155,252  
Filing Date: September 21, 1998  
Page 3 of 22

PATENT  
Attorney Docket No.: SALK1470-2  
(088802-1852)

*33*  
*Cont* from plasmid pSG424, described by Sadowski and Ptashne, in Nucleic Acids Res.  
17:7539 (1989) /--

Please replace the paragraph beginning at line 14 of page 23 with the following  
replacement paragraph:

*34* --pTK-PPRE3-LUC: Three copies of double-stranded peroxisome  
proliferator response element (PPRE) oligonucleotides (see SEQ ID NO:5) were  
cloned upstream of the TK promoter of TK-LUC at the *SaII* site. /--

Please amend the specification by entering the replacement Abstract of the Disclosure  
provided herein (a copy of the original page 40 of the specification as filed), enclosed as a  
separate sheet.

**In the claims:**

Please replace claims 16, 20, 27 and 28 with the following amended versions thereof: